Sr. Rights of No.   Description of rights and associated duty bearers	
1. Right to information  Every patient has a right to adequate relevant information about the nature, cause of illness, provisional/confirmed diagnosis, proposed investigations and management, and possible complications to be explained at their level of understanding in language  Every patient has a right to adequate hospital level 1 by Clinical Establishments set up as per Establishment Act 2010	
of illness, provisional/confirmed diagnosis, proposed investigations and management, and possible complications to be explained at their level of understanding in language	dard for
proposed investigations and management, and possible complications to be explained at their level of understanding in language	National
and possible complications to be explained at their level of understanding in language	Council
at their level of understanding in language	Clinical
known to them	
I who will be a horn.	
The treating physician has a duty to ensure 2) MCI Code of Ethics	
that this information is provided in simple	
and intelligible language to the patient to be 3) Patients Charter by	National
communicated either personally by the Accreditation Board for I	Hospitals
physician, or by means of his/her qualified (NABH)	
assistants.	
Every patient and his/her designated 4) The Consumer Protect	tion Act,
caretaker have the right to factual 1986	
information regarding the expected cost of	
treatment based on evidence. The hospital	
management has a duty to communicate this	
information in writing to the patient and	
his/her designated caretaker. They should	
also be informed about any additional cost to	
be incurred due to change in the physical	
condition of the patient or line of treatment in	
writing. On completion of treatment, The	
patient has the right to receive an itemized	
bill, to receive an explanation for the bill(s)	
regardless of the source of payment or the	
mode of payment, and receive payment	
receipt(s) for any payment made.	
Patients and their caretakers also have a	
right to know the identity and professional	
status of various care providers who are	
providing service to him/her and to know	

		which Doctor/Consultant is primarily	
		responsible for his/her care. The hospital	
		management has a duty to provide this	
		information routinely to all patients and their	
		caregivers in writing with an	
		acknowledgement.	
2.	Right to	Every patient or his caregiver has the right to 1)	Annexure 8 of standards for
	records and reports	access originals/copies of case papers,	hospital level 1 by National
	, opens	indoor patient records, investigation reports	Clinical Establishments Council
		(during period of admission, preferably within	set up as per clinical
		24 hours and after discharge, within 72	Establishment Act 2010
		hours) This may be made available wherever 2)	MCI Code of Ethics Section
		applicable after paying appropriate fees for	1.3.2
		photocopying allowed to be photocopied by 3)	Central Information Commission
		patients at their cost. The relatives/	Judgment, Nisha Priya Bhatia
		caregivers of the patient have to get	Vs. Institute of HB&AS, GNCTD,
		discharge summary or in case of death,	2014
		death summary along with original copies of 4)	) The Consumer protection Act.
		investigations. The hospital management	1986
		has a duty to provide these records and	
		reports and to instruct the responsible	
		hospital staff to ensure provision of the same	
		are strictly followed without fail.	
3	Right to	As per Supreme Court, all hospitals both in 1)	Supreme court judgment
	Emergency Medical Care	the government and in the private sector are	Parmanand katara v. Union of
		duty bound to provide basic Emergency	India (1989)
		Medical Care, and injured persons have a 2)	Judgment of National Consumer
		right to get Emergency Medical Care. Such	Disputes Redressal Commission
		care must be initiated without demanding	pravat kumar Mukherjee v. Ruby
		payment/advance and basic care should be	General Hospital & Others
		provided to the patient irrespective of paying	(2005)
		capacity. 3)	
		It is the duty of the hospital management to	and 2.4
		ensure provision of such emergency care 4	
		through its doctors and staff, rendered	'Right to Life'
		promptly without compromising on the	

		quality and safety of the patients.	
4	Right to	Every patient has a right that informed 1	) MCI code of Ethics section 7.76
	informed consent	consent must be sought prior to any 2	) Annexure 8 of standards for
	Consent	potentially hazardous test/treatment (e.g.	Hospital level 1 by National
		invasive investigation / surgery /	Clinical Establishments Council
		chemotherapy) which carries certain risks.	set up as per clinical
		It is the duty of the hospital management to	Establishment act 2010
		ensure that all concerned doctors are 3	) The Consumer Protection Act.
		properly instructed to seek informed consent	2010
		that an appropriate policy is adopted and 4	) Drugs and Cosmetic Act. 1940,
		that consent forms with protocol for seeking	Rules 2016 on Informed Consent
		informed consent, are provided for patients	
		in an obligatory manner.	
		It is the duty of the primary treating doctor	
		administering the potentially hazardous	
		test/treatment to explain to the patient and	
		caregivers the main risks that are involved in	
		the procedure, and after giving this	
		information, the doctor may proceed only if	
		consent has been given in writing by the	
		patient/caregiver or in the manner explained	
		under Drugs and cosmetic Act Rules 2016	
		on informed consent.	
5	Right to	All patients have a right to privacy, and 1	) MCI Code of Ethics sections
	confidentiality human dignity	doctors have a duty to hold information	2.2,7.14 and 7.17
	and privacy	about their health condition and treatment 2	) Annexure 8 of standards for
		plan in strict confidentiality, unless it is	Hospital level 1 by National
		essential in specific circumstances to	Clinical Establishments Council
		communicate such information in the interest	set up as per Clinical
		of protecting other or due to public health	Establishment Act 2010
		considerations.	
		Female patients have the right to presence	
		of another female person during physical	
		examination by a male practitioner; It is the	
		duty of the hospital management to ensure	
		presence of such female attendants in case	

		of female patients. The hospital		
		management has a duty to ensure that its		
		staff upholds the human dignity of every		
		patient in all situations. All data concerning		
		the patient should be kept under secured		
		safe custody and insulated from data theft		
		and leakage.		
6	Right to	Every patient has the right to seek second	1)	Annexure 8 of standards for
0	second		' /	
	opinion	opinion from an appropriate clinician of		Hospital level 1 by National
		patients/caregivers choice. The hospital		Clinical Establishments Council
		management has a duty to respect the		set up as per Clinical
		patient's right to second opinion, and should	٥,	Establishment Act 2010
		provide to the patients caregivers all	2)	The Consumer Protection Act,
		necessary records and information required		1986
		for seeking such opinion without any extra		
		cost or delay.		
		The hospital management has a duty to		
		ensure that any decision to seek such		
		second opinion by the patient/caregivers		
		must not adversely influence the quality of		
		care being provided by the treating hospital		
		as long as the patient is under care of that		
		hospital. Any kind discriminatory practice		
		adopted by the hospital or the service		
		providers will be deemed as Human Rights		
		violation.		
7	Right to	Every patient and their caregivers have a	1)	MCI Code of Ethics Section 1.8
	transparency in rates, and	right to information on the rates to be		regarding Payment of
	care	charged by the hospital for each type of		Professional Services
	according to prescribed	service provided and facilities available on a	2)	Section 9(i) and 9(ii) of Clinical
	rates	prominent display board and a brochure.		establishments (Central
	wherever relevant	They have a right to receive a itemized		Government) Rules 2012
	reievant	detailed bill at the time of payment. It would	3)	Annexure 8 of standards for
		be the duty of hospital/Clinical Establishment		hospital level 1 by National
		to display key rates at a conspicuous place		Clinical Establishment Council
		in local as well as English language, and to		set up as per clinical
				· '

		make available the detailed schedule of		Establishment Act 2010
		rates in a booklet form to all	4)	Various Drug price control
		patients/caregivers.		orders
		Every patient has a right to obtain essential	5)	The Consumer Protection Act.
		medicines as per India pharmacopeia,		1986
		devices Authority (NPPA) and other relevant	6)	Drugs price control Order
		authorities. Every patient has a right to		(DPCO) section 3 of the
		receive health care services within the range		Essential Commodities Act,
		of rates for procedures and services		1955
		prescribed by central and state Governments		
		from time to time, wherever relevant.		
		However, no patient can be denied choice in		
		terms of medicines, devices and standard		
		treatment guidelines based on the		
		affordability of the patients' right to choice.		
		Every hospital and clinical establishment has		
		a duty to ensure that essential medicines		
		under NLEM as per Government of India and		
		World Health Organization, devices, implants		
		and services are provided to patients at rates		
		that are not higher than the prescribed rates		
		or the maximum retail price marked on the		
		packaging.		
8	Right to non- discrimination	Every patient has the right to receive	1)	Annexure 8 of standards for
	discrimination	treatment without any discrimination based		Hospital level 1 by National
		on his or her illnesses or conditions,		Clinical Establishments Council
		including HIV status or other health		set up as per Clinical
		condition, religion, caste, ethnicity, gender,		Establishment Act 2010
		age, sexual orientation, linguistic or		
		geographical/social origins.		
		The hospital management has a duty to		
		ensure that no form of discriminatory		
		behavior or treatment takes place with any		

person under the hospital's care. The hospital management must regularly orient and instruct all its doctors and staff regarding

		the same.		
9	Right to	Patients have a right to safety and security in	1)	Clinical establishments (Central
	safety and quality care	the hospital premises. They have a right to		Government) Rules 2012
	according to standards	be provided with care in an environment	2)	The Consumer Protection Act.
		having requisite cleanliness, infection control		1986
		measures, safe drinking water as per		
		BIS/FSSAI Standards and sanitation		
		facilities. The hospital management has a		
		duty to ensure safety of all patients in its		
		premises including clean premises and		
		provision for infection control. Patients have		
		a right to receive quality health care		
		according to currently accepted standards,		
		norms and standard guidelines as per		
		National Accreditation Board for Hospitals		
		(NABH) or similar. They have a right to be		
		attended to, treated and cared for with due		
		skill, and in a professional manner in		
		complete consonance with the principles of		
		medical ethics. Patients and caretakers have		
		a right to seek redressal in case of perceived		
		medical negligence or damaged caused due		
		to deliberate deficiency in service delivery.		
		The hospital Management and treating		
		doctors have a duty to provide quality health		
		care in accordance with current standards of		
		care and standard treatment guidelines and		
		to avoid medical negligence or deficiency in		
		service delivery system in any form.		
10	Right to	Patients and their caregivers have a right to	1)	Annexure 8 of standards for
	choose alternative	choose between alternative treatment/		Hospital level 1 by National
	treatment options if	management options, if these are available,		Clinical Establishments Council
	options if available	after considering all aspects of the situation.		set up as per Clinical
		This includes the option of the patient		Establishment Act 2010
		refusing care after considering all available	2)	The Consumer Protection Act,
		options, with responsibility for consequences		1986
1				

		being borne by the patient and his/her	
		caregivers. In case a patient leaves a	
		healthcare facility against medical advice on	
		his/her own responsibility, then	
		notwithstanding the impact that this may	
		have on the patient's further treatment and	
		condition, this decision itself should not	
		affect the observance of various rights	
		mentioned in this charter.	
		The hospital management has a duty to	
		provide information about such options to the	
		patient as well as to respect the informed	
		choice of the patient and caregivers in a	
		proper recorded manner with due	
		acknowledgement from the patient or the	
		caregivers on the communication and the	
		mode.	
11	Right to	When any medicine is prescribed by a doctor	1) Various judgments by the
11	Right to choose source for	When any medicine is prescribed by a doctor or a hospital, the patients and their	Various judgments by the     National Consumer Dispute
11	choose source for obtaining	·	
11	choose source for	or a hospital, the patients and their	National Consumer Dispute
11	choose source for obtaining medicines or	or a hospital, the patients and their caregivers have the right to choose any	National Consumer Dispute Redressal Commission
11	choose source for obtaining medicines or	or a hospital, the patients and their caregivers have the right to choose any registered pharmacy of their choice to	National Consumer Dispute Redressal Commission 2) The Consumer Protection
11	choose source for obtaining medicines or	or a hospital, the patients and their caregivers have the right to choose any registered pharmacy of their choice to purchase them. Similarly when a particular	National Consumer Dispute Redressal Commission 2) The Consumer Protection
11	choose source for obtaining medicines or	or a hospital, the patients and their caregivers have the right to choose any registered pharmacy of their choice to purchase them. Similarly when a particular investigation is advised by a doctor or a	National Consumer Dispute Redressal Commission 2) The Consumer Protection
11	choose source for obtaining medicines or	or a hospital, the patients and their caregivers have the right to choose any registered pharmacy of their choice to purchase them. Similarly when a particular investigation is advised by a doctor or a hospital, the patient and his caregiver have a	National Consumer Dispute Redressal Commission 2) The Consumer Protection
11	choose source for obtaining medicines or	or a hospital, the patients and their caregivers have the right to choose any registered pharmacy of their choice to purchase them. Similarly when a particular investigation is advised by a doctor or a hospital, the patient and his caregiver have a right to obtain this investigation from any	National Consumer Dispute Redressal Commission 2) The Consumer Protection
11	choose source for obtaining medicines or	or a hospital, the patients and their caregivers have the right to choose any registered pharmacy of their choice to purchase them. Similarly when a particular investigation is advised by a doctor or a hospital, the patient and his caregiver have a right to obtain this investigation from any registered diagnostic centre/laboratory	National Consumer Dispute Redressal Commission 2) The Consumer Protection
11	choose source for obtaining medicines or	or a hospital, the patients and their caregivers have the right to choose any registered pharmacy of their choice to purchase them. Similarly when a particular investigation is advised by a doctor or a hospital, the patient and his caregiver have a right to obtain this investigation from any registered diagnostic centre/laboratory having qualified personnel and accredited by	National Consumer Dispute Redressal Commission 2) The Consumer Protection
11	choose source for obtaining medicines or	or a hospital, the patients and their caregivers have the right to choose any registered pharmacy of their choice to purchase them. Similarly when a particular investigation is advised by a doctor or a hospital, the patient and his caregiver have a right to obtain this investigation from any registered diagnostic centre/laboratory having qualified personnel and accredited by National Accreditation Board for Laboratories (NABL).  It is the duty of every treating	National Consumer Dispute Redressal Commission 2) The Consumer Protection
11	choose source for obtaining medicines or	or a hospital, the patients and their caregivers have the right to choose any registered pharmacy of their choice to purchase them. Similarly when a particular investigation is advised by a doctor or a hospital, the patient and his caregiver have a right to obtain this investigation from any registered diagnostic centre/laboratory having qualified personnel and accredited by National Accreditation Board for Laboratories (NABL).	National Consumer Dispute Redressal Commission 2) The Consumer Protection
11	choose source for obtaining medicines or	or a hospital, the patients and their caregivers have the right to choose any registered pharmacy of their choice to purchase them. Similarly when a particular investigation is advised by a doctor or a hospital, the patient and his caregiver have a right to obtain this investigation from any registered diagnostic centre/laboratory having qualified personnel and accredited by National Accreditation Board for Laboratories (NABL).  It is the duty of every treating physician/hospital management to inform the patient and his caregivers that they are free	National Consumer Dispute Redressal Commission 2) The Consumer Protection
11	choose source for obtaining medicines or	or a hospital, the patients and their caregivers have the right to choose any registered pharmacy of their choice to purchase them. Similarly when a particular investigation is advised by a doctor or a hospital, the patient and his caregiver have a right to obtain this investigation from any registered diagnostic centre/laboratory having qualified personnel and accredited by National Accreditation Board for Laboratories (NABL).  It is the duty of every treating physician/hospital management to inform the	National Consumer Dispute Redressal Commission 2) The Consumer Protection
11	choose source for obtaining medicines or	or a hospital, the patients and their caregivers have the right to choose any registered pharmacy of their choice to purchase them. Similarly when a particular investigation is advised by a doctor or a hospital, the patient and his caregiver have a right to obtain this investigation from any registered diagnostic centre/laboratory having qualified personnel and accredited by National Accreditation Board for Laboratories (NABL).  It is the duty of every treating physician/hospital management to inform the patient and his caregivers that they are free	National Consumer Dispute Redressal Commission 2) The Consumer Protection

patient/ caregiver to access pharmacy/

		diagnostic centre of their choice must not in	
		any ways adversely influence the care being	
		provided by the treating physician or	
		hospital.	
12	Right to	A patient has the right to continuity of care,	1) Medical Council of India
	proper referral and	and the right to be duly registered at the first	code of ethics section 3.6
	transfer,	healthcare facility where treatment has been	2) World Health Organization
	which is free from perverse	sought, as well as at any subsequent	Referral Notes
	commercial	facilities where care is sought, when being	3) Various IPHS documents
	influences	transferred from one healthcare facility to	
		another, the patient/caregiver must receive a	
		complete explanation of the justification for	
		the transfer, the alternative options for a	
		transfer and it must be confirmed that the	
		transfer is acceptable to the receiving facility.	
		The patient and caregivers have right to be	
		informed by the hospital about any	
		continuing healthcare requirements following	
		discharge from the hospital. The hospital	
		management has a duty to ensure proper	
		referral transfer of patients regarding such a	
		shift in care.	
		In regard to all referrals of patients, including	
		referrals to other hospitals, specialists,	
		laboratories or imaging services, the	
		decision regarding facility to which referral is	
		made must be guided entirely by the best	
		interest of the patient. The referral process	
		must not be influenced by any commercial	
		consideration such as kickbacks,	
		commissions, incentives, or other perverse	
		business practices.	
13	Right to	Every person/patient who is approached to	1) Protocols and Good Clinical
	protection for patients	participate in a clinical trial has a right to due	Practice Guidelines issued by
	involved in	protection in this context. All clinical trials	Central Drugs Standard control
	clinical trials	must be conducted in compliance with the	organization, Directorate

protocols and clinical practice good guidelines issued by central Drugs Standard control Organization, Directorate General of 2 Amended Drugs and Cosmetics Health Services, Govt. of India as well as all applicable statutory provisions of Amended Drugs and Cosmetics Act, 1940 and Rules, 3) 1945, including observance of the following provisions related to patients rights:

- a) Participation of patients in clinical trials must always be based on informed consent, given after provision of all 4) World relevant information. The patient must be given a copy of the signed informed consent form, which provides him/ her with a record containing basic information about the trial and also becomes documentary evidence to prove their participation in the trial.
- b) A participant's right to agree or decline consent to take part in a clinical trial must be respected and her/ his refusal should not affect routine care.
- c) The patient should also be informed in writing about the name of the drug/ intervention that is undergoing trial along with dates. dose and duration administration.
- d) At all times, the privacy of a trial participant must be maintained and any information gathered from the participant must be kept strictly confidential.
- e) Trial participants who suffer any adverse impact during their participation in a trial are entitled to free medical management events, adverse irrespective relatedness to the clinical trial, which

- General of Health Services. Govt. of India
- Act, 1940 and Rules, 1945 especially schedule Y
- National Ethical Guidelines for Biomedical and Health Research involving Human Participants, Indian Council of Medical Research New Delhi, 2017
- Medical Assembly Declaration of Helsinki: Ethical Principles for Medical Research Involvina Human Subjects available at

www.wma.net/en/30publications/ 10policies/b3/17c.pdf

		participants should follow the national Ethical 2) World Medical Assembly Guidelines for Biomedical and Health Declaration of Helsinki: Ethical
	and health research	participant has a right to due protection in this context. Any research involving such Research, New Delhi, 2017
	involved in biomedical	research participant and every research Involving Human Participants,
	participants	biomedical research shall be referred to as Biomedical and Health Research
14	Right to protection of	Every patient who is taking part in 1) National Ethical Guidelines for
4.4	Diabéta	persons/ patients involved in such a trial.
		these guidelines are followed in case of any
		clinical trial has a duty to ensure that all
		Any doctor or hospital who is involved in a
		study.
		methods that my have been proven by the
		assured of access to the best treatment
		h) After the trial, participants should be
		Committee.
		necessary by the concerned Ethics
		compensation wherever deemed
		(ancillary care) and award of
		of trial related or unrelated illnesses
		g) Institutional mechanisms must be established to allow insurance coverage
		appropriate.  g) Institutional mechanisms must be
		care or reference to facilities, as may be
		trial. This could be in the form of medical
		illnesses arising during the period of the
		trial participants for non-study/trial related
		f) Ancillary care may be provided to clinical
		compensation.
		death, their dependents have the right to
		any impairment or disability. In case of
		must be given to compensate them for
		addition, financial or other assistance
		injury is not related to the clinical trial. In
		till such time as it is established that the
		should be given for as long as required or

Research and should be carried out with prior approval of the Ethics Committee.  Documented informed consent of the 10policies/b3/17c.pdf			Research involving Human Participants,	Principles for Medical Research
prior approval of the Ethics Committee.  Documented informed consent of the research participants should be taken.  Additional safeguards should be taken in research involving vulnerable population.  Right to privacy and confidentiality  Research participants who suffer any direct psychological, social, legal or economic harm as a result of their participation are entitled, after due assessment, to financial or other assistance to compensate them equitably for any temporary or permanent			2017 laid down by Indian council for Medical	Involving Human Subjects
Documented informed consent of the research participants should be taken. Additional safeguards should be taken in research involving vulnerable population. Right to privacy and confidentiality Research participants who suffer any direct psychological, social, legal or economic harm as a result of their participation are entitled, after due assessment, to financial or other assistance to compensate them equitably for any temporary or permanent			Research and should be carried out with	available at
research participants should be taken. Additional safeguards should be taken in research involving vulnerable population. Right to privacy and confidentiality Research participants who suffer any direct psychological, social, legal or economic harm as a result of their participation are entitled, after due assessment, to financial or other assistance to compensate them equitably for any temporary or permanent			prior approval of the Ethics Committee.	www.wma.net/en/30publications/
Additional safeguards should be taken in research involving vulnerable population. Right to privacy and confidentiality Research participants who suffer any direct psychological, social, legal or economic harm as a result of their participation are entitled, after due assessment, to financial or other assistance to compensate them equitably for any temporary or permanent			Documented informed consent of the	10policies/b3/17c.pdf
research involving vulnerable population. Right to privacy and confidentiality Research participants who suffer any direct psychological, social, legal or economic harm as a result of their participation are entitled, after due assessment, to financial or other assistance to compensate them equitably for any temporary or permanent			research participants should be taken.	3) Drugs & Cosmetic Act. Rules
Right to privacy and confidentiality Research participants who suffer any direct psychological, social, legal or economic harm as a result of their participation are entitled, after due assessment, to financial or other assistance to compensate them equitably for any temporary or permanent			Additional safeguards should be taken in	2016 on Clinical Trails
Research participants who suffer any direct psychological, social, legal or economic harm as a result of their participation are entitled, after due assessment, to financial or other assistance to compensate them equitably for any temporary or permanent			research involving vulnerable population.	
psychological, social, legal or economic harm as a result of their participation are entitled, after due assessment, to financial or other assistance to compensate them equitably for any temporary or permanent			Right to privacy and confidentiality	
harm as a result of their participation are entitled, after due assessment, to financial or other assistance to compensate them equitably for any temporary or permanent			Research participants who suffer any direct	
entitled, after due assessment, to financial or other assistance to compensate them equitably for any temporary or permanent			psychological, social, legal or economic	
other assistance to compensate them equitably for any temporary or permanent			harm as a result of their participation are	
equitably for any temporary or permanent			entitled, after due assessment, to financial or	
			other assistance to compensate them	
impairment or disability.			equitably for any temporary or permanent	
			impairment or disability.	
The benefits accruing from research should			The benefits accruing from research should	
be made accessible to individuals,			be made accessible to individuals,	
communities and populations whenever			communities and populations whenever	
relevant. Any doctor or hospital who is			relevant. Any doctor or hospital who is	
involved in biomedical and health research			involved in biomedical and health research	
involving patients has a duty to ensure that			involving patients has a duty to ensure that	
all these guidelines are followed in case of			all these guidelines are followed in case of	
any persons/ patients involved in such			any persons/ patients involved in such	
research.			research.	
	15		A patient has the right to take discharge and	1) Prohibition of wrongful
discharge of patient, or cannot be detained in a hospital on confinement under sec. 340-34			cannot be detained in a hospital on	confinement under sec. 340-342
receive body procedural grounds such as dispute in of IPC. Statements of Mumb			procedural grounds such as dispute in	of IPC. Statements of Mumbai
from hospital payment of hospital charges. Similarly, High Court.			payment of hospital charges. Similarly,	High Court.
caretakers have the right to the dead body of 2) Consumer Protection Act. 1986		-	caretakers have the right to the dead body of	2) Consumer Protection Act. 1986
a patient who had been treated in a hospital			a patient who had been treated in a hospital	
and the dead body cannot be detailed on			and the dead body cannot be detailed on	
procedural grounds. Including nonpayment/			procedural grounds. Including nonpayment/	
dispute regarding payment of hospital			l l	
charges against wishes of the caretakers.			dispute regarding payment of hospital	
The hospital management has a duty to				

	T			
		observe these rights and not to indulge in		
		wrongful confinement of any patient, or dead		
		body of patient, treated in the hospital under		
		any circumstances.		
16	Right to Patient	Patients have the right to receive education	1)	The Consumer Protection Act,
	Education	about major facts relevant to his/her		1986
		condition and healthy living practices, their	2)	Standards for Hospital level 1 by
		rights and responsibilities, officially		National Clinical Establishments
		supported heath insurance schemes relevant		Council set up as per Clinical
		to the patient, relevant entitlements' in case		Establishment Act. 2010
		of charitable hospitals, and how to seek		
		redressal of grievances in the patients		
		understand or seek the education.		
		The hospital management and treating		
		physician have a duty to provide such		
		education to each patient according to		
		standard procedure in the language the		
		patients understand and communicate in a		
		simple and easy to understand manner.		
17	Right to be	Every patient and their caregivers have the	1)	The Consumer Protection Act,
	heard and seek	right to give feedback, make comments, or		1986
	redressal	lodge complaints about the health care they	2)	NHS-Charter of Patient Rights
		are receiving or had received from a doctor		and Responsibilities
		or hospital. This includes the right to be		
		given information and advice on how to give		
		feedback, make comments, or make a		
		complaint in a simple and user friendly		
		manner.		
		Patients and caregivers have the right to		
		seek redressal in case they are aggrieved,		
		on account of infringement of any of the		
		above mentioned rights in this charter. This		
		may be done by lodging a complaint with an		
		official designated for this purpose by the		
		hospital/healthcare provider and further with		
		an official mechanism constituted by the		

government such as patients' rights Tribunal Forum or clinical establishments regulatory authority as the case may be. All complaints must be registered by providing a registration number and there should be a robust tracking and tracing mechanism to ascertain the status of the complaint resolution.

The patient and caregivers have the right to a fair and prompt redressal of their grievances. Further, they have the right to receive in writing the outcome of the complaint within 15 days from the date of the receipt of the complaint.

Every hospital and clinical establishment has the duty to set up an internal redressal mechanism as well as to fully comply and cooperate with official redressal mechanisms including making available all relevant information and taking action in full accordance with orders of the redressal body as per the patient's Right charter or as per the applicable existing laws.